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**INFORMATION DISCLOSURE
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(Not for submission under 37 CFR 1.99)

Application Number	10722374
Filing Date	2003-11-25
First Named Inventor	David Bebbington
Art Unit	1624
Examiner Name	Venkataraman Balasubramanian
Attorney Docket Number	VPI/00-130-08 CON US

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	1	Tanzi, K. et al., "Purines. X. Reactivities of Methyl Groups on 9-Phenylpurines : Condensation with an Aldehyde or an Ester, and Oxidation with Selenium Dioxide", <i>Chem. Phar. Bull.</i> , 40 (1), 227-229 (1992).	<input type="checkbox"/>
	2	Charpiot, B. et al., "Quinazolines: Combined type 3 and 4 phosphodiesterase inhibitors", <i>Bioorg. Med. Chem. Lett.</i> , 8 (20), 2891-2896 (1998).	<input type="checkbox"/>
	3	Shikhaliev, K.S. et al., "Heterocyclization of quinazol-2-ylguanidines. 1. Reaction with amino acids", <i>Chem. Heterocycl. Compd.</i> , 35 (7), 818-820 (1999).	<input type="checkbox"/>
	4	Singh, S.P. et al., "Synthesis & Mass Spectra of Some Substituted 2-(2'-Benzazolylamino)pyrimidines", <i>Indian J. Chem. Sect. B</i> , 22(1); 37-42 (1983).	<input type="checkbox"/>
	5	Ti, J. et al., "Anticandidal activity of pyrimidine-peptide conjugates", <i>J. Med. Chem.</i> , 23(8), 913 – 918 (1980).	<input type="checkbox"/>
	6	Kretzschmar, E. et al., "Synthese von 2,6-disubstituierten 4-Hydroxy-5,6,7,8-tetrahydropyrido[4,3-d]pyrimidinen", <i>Pharmazie</i> , 43(7), 475-476 (1988).	<input type="checkbox"/>
	7	Norman, M.H. et al., "Structure-Activity Relationships of a Series of Pyrrolo[3,2-d]pyrimidine Derivatives and Related Compounds as Neuropeptide Y5 Receptor Antagonists", <i>J. Med. Chem.</i> , 43(22), 4288 -4312 (2000).	<input type="checkbox"/>
	8	Nugent, R.A. et al., "Pyrimidine Thioethers: A Novel Class of HIV-1 Reverse Transcriptase Inhibitors with Activity Against BHAP-Resistant HIV", <i>J. Med. Chem.</i> , 41, 3793-3803 (1998).	<input type="checkbox"/>
	9	Myers, M.R. et al., "The synthesis and SAR of new 4-(N-alkyl-N-phenyl)amino-6,7-dimethoxyquinazolines and 4-(N-alkyl-N-phenyl)aminopyrazolo[3,4-d]pyrimidines, inhibitors of CSF-1R tyrosine kinase activity", <i>Bioorg. Med. Chem. Lett.</i> , 7, 4, 421-424 (1997).	<input type="checkbox"/>
	10	Agarwal, N. et al., "Suitably functionalised pyrimidines as potential antimycotic agents", <i>Bioorg. Med. Chem. Lett.</i> , 10, 8, 703-706 (2000).	<input type="checkbox"/>
	11	Crespo, M.I. et al., "Design, Synthesis, and Biological Activities of New Thieno[3,2-d]pyrimidines as Selective Type 4 Phosphodiesterase Inhibitors", <i>J. Med. Chem.</i> , 41 (21), 4021 -4035 (1998).	<input type="checkbox"/>

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	12	Noell, C.W. et al., "Potential Purine Antagonists. XX. The Preparation and Reactions of Some Methylthiopurines", <i>J. Am. Chem. Soc.</i> , 81(22), 5997 – 6007 (1959).	<input type="checkbox"/>
	13	Lubbers, T. et al., "Design, synthesis, and structure–activity relationship studies of ATP analogues as DNA gyrase inhibitors", <i>Bioorg. Med. Chem. Lett.</i> , 10, 8, 821-826 (2000).	<input type="checkbox"/>
	14	D'Atri, G. et al., "Novel pyrimidine and 1,3,5-triazine hypolipemic agents", <i>J. Med. Chem.</i> 27(12), 1621 – 1629 (1984).	<input type="checkbox"/>
	15	Venugopalan, B. et al., "Synthesis and antimalarial activity of pyrido[3,2-f]quinoxalines and their oxides", <i>Indian J. Chem. Sect. B</i> , 34, 9, 778-790 (1995).	<input type="checkbox"/>
	16	Curd, F.H.S. et al, "Synthetic antimalarials. Part XVII. Some aminoalkylaminoquinoline derivatives", <i>J. Chem. Soc.</i> , 899 – 909 (1947).	<input type="checkbox"/>
	17	Haworth, R.D. et al., "Synthetic antimalarials. Part XXVII. Some derivatives of phthalazine, quinoxaline, and isoquinoline", <i>J. Chem. Soc.</i> , 777 – 782 (1948).	<input type="checkbox"/>
	18	Nair, M.D., et al., "3-Chloroisocarbostyryl & Its Chlorination Products", <i>Indian J. Chem.</i> , 467-470 (1967).	<input type="checkbox"/>
	19	Jeffery, J.E. et al., "Synthesis of sibutramine, a novel cyclobutylalkylamine useful in the treatment of obesity, and its major human metabolites", <i>J. Chem. Soc., Perkin Trans. 1</i> , 21, 2583-2589 (1996).	<input type="checkbox"/>
	20	Cohen, P., "Dissection of the Protein Phosphorylation Cascades Involved in Insulin and Growth Factor Action", <i>Biochem. Soc. Trans.</i> , 21, 555-567 (1993).	<input type="checkbox"/>
	21	Haq, S. et al., "Glycogen Synthase Kinase-3 β Is a Negative Regulator of Cardiomyocyte Hypertrophy", <i>J. Cell Biol.</i> , 151(1), 117-129 (2000).	<input type="checkbox"/>
	22	Fischer, P.M. et al., "Inhibitors of Cyclin-Dependent Kinases as Anti-Cancer Therapeutics", <i>Current Med. Chem.</i> , 7, 1213-1245 (2000).	<input type="checkbox"/>

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	23	Mani, S. et al., "Cyclin-dependent kinase: novel anticancer agents", <i>Exp. Opin. Invest. Drugs.</i> , 8, 1849-1870 (2000). <input type="checkbox"/>
	24	Fry, D.W. et al., "Inhibitors of cyclin-dependent kinases as therapeutic agents for the treatment of cancer", <i>Current Opin. Oncol. Endoc. & Metab. Investig.</i> , 2-40-59 (2000). <input type="checkbox"/>
	25	Bokemeier, D. et al., "Multiple intracellular MAP kinase signaling cascades", <i>Kidney Int.</i> , 49, 1187-1198 (1996). <input type="checkbox"/>
	26	Anderson, N.G. et al., "Multiple intracellular MAP kinase signaling cascades", <i>Nature</i> , 343, 651-653 (1990). <input type="checkbox"/>
	27	Crews, C.M. et al., "The Primary Structure of MEK, a Protein Kinase That Phosphorylates the ERK Gene Product", <i>Science</i> , 258, 478-480 (1992). <input type="checkbox"/>
	28	Bjorbaek, C. et al, "Divergent Functional Roles for p90rsk Kinase Domains", <i>J. Biol. Chem.</i> , 270(32), 18848-18552 (1995). <input type="checkbox"/>
	29	Rouse, J. et al., A Novel Kinase Cascade Triggered by Stress and Heat Shock That Stimulates MAPKAP Kinase-2 and Phosphorylation of the Small Heat Shock Proteins", <i>Cell</i> , 78, 1027-1037 (1994). <input type="checkbox"/>
	30	Raingeaud, J. et al., MMK3- and MMK6-Regulated Gene Expression Is Mediated by p38 Mitogen-Activated Protein Kinase Signal Transduction Pathway", <i>Mol. Cell. Biol.</i> , 16, 1247-1255 (1996). <input type="checkbox"/>
	31	Chen, R.H. et al., "Phosphorylation of the c-Fos transrepression domain by mitogen-activated protein kinase and 90-kDa ribosomal S6 kinase", <i>Proc. Natl. Acad. Sci. USA</i> , 90, 10952-10956 (1993). <input type="checkbox"/>
	32	Moodie, S.A. et al., "Complexes of Ras-GTP with Raf-1 and Mitogen-Activated Protein Kinase Kinase", <i>Science</i> , 260 (5114), 1658-1661 (1993). <input type="checkbox"/>
	33	Frey, R.S. et al., "Involvement of Extracellular Signal-regulated Kinase 2 and Stress-activated Protein Kinase/Jun N-Terminal Kinase Activation by Transforming Growth Factor β in the Negative Growth Control of Breast Cancer Cells", <i>Cancer Res.</i> , 57, 628-633 (1997). <input type="checkbox"/>

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	34	Sivaraman, V.S., et al., "Hyperexpression of Mitogen-activated Protein Kinase in Human Breast Cancer", J. Clin. Invest., 99(7), 1478-1483 (1997). <input type="checkbox"/>
	35	Whelchel, A. et al., "Inhibition of ERK Activation Attenuates Endothelin-stimulated Airway Smooth Muscle Cell Proliferation", Am. J. Respir. Cell Mol. Biol., 16, 589-596 (1997). <input type="checkbox"/>
	36	Yuan, Z.Q. et al., "Frequent activation of AKT2 and induction of apoptosis by inhibition of phosphoinositide-3-OH kinase/Akt pathway in human ovarian cancer", Oncogene, 19, 2324-2330 (2000). <input type="checkbox"/>
	37	Kazuhiko, N. et al., "Akt/Protein Kinase B Prevents Injury-Induced Motoneuron Death and Accelerates Axonal Regeneration", J. of Neuroscience, 20(8), 2875-2986 (2000). <input type="checkbox"/>
	38	Molina, T.J. et al., "Profound block in thymocyte development in mice lacking p56lck", Nature, 357, 161-164 (1992). <input type="checkbox"/>
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